Amendments to the Claims:

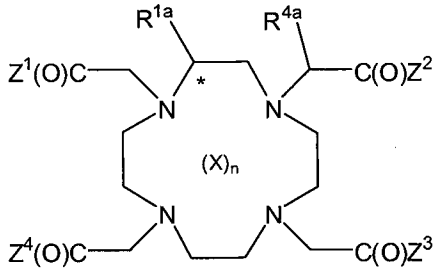
This listing of claims will replace all prior versions, and listings of claims in the application.

Listing of Claims:

1. (Canceled).

2. – 5. (Canceled).

6. (Previously presented) The method of claim 43, wherein said substituted or unsubstituted 1,4,7,10-tetraazacyclododecane-N,N',N",N"'-tetraacetic acid (DOTA) has the formula:



wherein

R^{1a} and R^{4a} are members independently selected from H, substituted or unsubstituted alkyl, substituted or unsubstituted heteroalkyl, substituted or unsubstituted aryl and linker moieties;

X is a member selected from a lanthanide ion, an actinide ion, an alkaline earth metal ion, and a group IIIb transition metal ion;

Z^1, Z^2, Z^3 and Z^4 are members independently selected from OR^1 and NR^1R^2

in which

R^1 and R^2 are members independently selected from H, substituted or unsubstituted alkyl and substituted or unsubstituted heteroalkyl;

n is a member selected from 0 and 1.

7. (Cancelled).

8. (Previously presented) The method of claim **6**, wherein the carbon atom marked * is of S configuration.

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9. (Cancelled)

10. (Previously presented) The method of claim **43**, wherein said targeting moiety binds specifically to said cell surface antigen.

11. (Previously presented) The method of claim **43**, wherein the targeting moiety is covalently attached to said antibody.

12. (Previously presented) The method of claim **10**, wherein the targeting moiety is a second antibody.

13. (Original) The method of claim **11**, wherein the targeting moiety specifically binds to a protein on a cancer cell.

14. (Previously presented) The method of claim **43**, wherein the subject is a mammal.

15. (Previously presented) The method of claim **14**, wherein the mammal is a human.

16-23. (Canceled)

24. (Previously presented) The method according to claim **43** wherein said antibody has the structure:

$$(Ab)_{n'}\text{-}L\text{-}T$$

wherein,

n′ is an integer selected from 1 to 10 ;

Ab represents said antibody;

L is a member selected from a chemical bond and a linking group that may contain one or more functional groups; and

T is said targeting moiety.

25. (Canceled).

26. (Previously presented) The method of claim **24**, wherein said targeting moiety is a second antibody that binds specifically to a cell surface antigen.

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27. (Previously presented) The method according to claim **24** wherein said antibody is administered to said subject as a pharmaceutical composition comprising said antibody and a pharmaceutically acceptable carrier.

28. (Canceled)

29. (Cancelled).

30. (Canceled)

31 (Canceled)

32. (Cancelled).

33. (Previously presented) The method according to claim **6**, wherein

R^{1a} and R^{4a} are H;

Z^1, Z^2, Z^3 and Z^4 are OH;

and n is 1.

34. (Previously presented) The method according to claim **33**, wherein said targeting moiety is a second antibody that binds specifically to a cell surface antigen.

35. (Previously presented) The method according to claim **34**, wherein said targeting moiety is anti-CEA.

36. (Previously presented) The method according to claim **33**, wherein said targeting moiety is anti-CEA.

37. (Canceled)

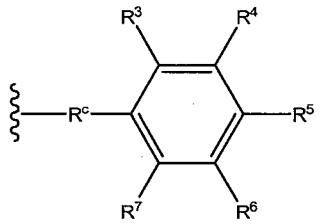
38. (Previously presented) The method according to claim **43**, wherein said reactive site comprises sulfur.

39. (Cancelled)

40. (Previously presented) The method according to claim **6**, wherein R^{1a} is a member independently selected from substituted or unsubstituted alkyl, substituted or unsubstituted heteroalkyl, substituted or unsubstituted aryl and linker moieties.

41. (Previously presented) The method according to claim **6**, wherein R^{4a} is a member independently selected from substituted or unsubstituted alkyl, substituted or unsubstituted heteroalkyl, substituted or unsubstituted aryl and linker moieties.

42. (Previously presented) The method according to claim **6**, wherein said DOTA further comprises an arylalkyl moiety having a structure according to the formula:



wherein

R^c is an unsubstituted unbranched alkyl linker;

R^3, R^4, R^5, R^6 and R^7 are members independently selected from H, halogen, NO_2, CN, X^1R^8, NR^9R^{10}, and $C(X^2)R^{11}$,

wherein

X^1 is a member selected from O, NH, and S;

X^2 is a member selected from O, S, and NH;

R^8 and R^9 are members independently selected from H, substituted or unsubstituted alkyl, substituted or unsubstituted heteroalkyl, substituted or unsubstituted heteroalkyl and $C(Z^3)R^{12}$

wherein

Z^3 is a member selected from O, S and NH;

R^{12} is a member selected from substituted or unsubstituted alkyl, substituted or unsubstituted heteroalkyl and OR^{13}

wherein

R^{13} is a member selected from substituted or unsubstituted alkyl, substituted or unsubstituted heteroalkyl, substituted or unsubstituted aryl, and substituted or unsubstituted heteroaryl;

R^{10} is a member selected from H, substituted or unsubstituted alkyl, substituted or unsubstituted heteroalkyl, and OH, and

R^9 and R^{10} taken together are optionally (=C=S);

R^{11} is a member selected from H, halogen, substituted or unsubstituted alkyl, substituted or unsubstituted heteroalkyl, OR^{14}, and $NR^{15}R^{16}$,

wherein

R^{14} is a member selected from H, substituted or unsubstituted alkyl, substituted or unsubstituted heteroalkyl, and $C(O)R^{17}$,

wherein

R^{17} is a member selected from substituted or unsubstituted alkyl, and substituted or unsubstituted heteroalkyl; and

R^{15} and R^{16} are members independently selected from H, substituted or unsubstituted alkyl, and substituted or unsubstituted heteroalkyl

43. (Previously presented) A method of treating a subject with cancer by administration of a macrocyclic metal chelate, said method comprising the steps of:

(a) administering to said subject an antibody comprising an antigen recognition domain that recognizes said macrocyclic metal chelate, wherein said antibody comprises:

i) a light chain comprising:

a) a first CDR having the sequence of SEQ ID NO:2;

b) a second CDR having a sequence selected from the group consisting of:

i) SEQ ID NO:3; and

ii) SEQ ID NO:3 containing a cysteine substitution wherein position 2 is substituted by a cysteine;

c) a third CDR having the sequence of SEQ ID NO:4;

ii) a heavy chain comprising:

a) a first CDR having the sequence of SEQ ID NO:6;

b) a second CDR having a sequence selected from the group consisting of:

i) SEQ ID NO:7;

ii) SEQ ID NO: 7 containing a cysteine substitution wherein position 5 has been substituted by a cysteine;

iii) SEQ ID NO:7 containing a cysteine substitution wherein position 6 has been substituted by a cysteine; and

iv) SEQ ID NO:7 containing a cysteine substitution wherein position 7 has been substituted by a cysteine;

c) a third CDR having the sequence of SEQ ID NO:8; wherein said antibody comprises at least one of said cysteine substitutions, and wherein said antibody binds substituted or unsubstituted 1,4,7,10-tetraazacyclododecane-N,N',N",N"'-tetraacetic acid (DOTA); and

a targeting moiety that binds specifically to a cancer cell by binding with a member selected from a cell surface receptor and cell surface antigen, thereby forming a cell-antibody complex; and

(b) administering to said subject said macrocyclic metal chelate, thereby forming a covalent bond between said reactive site and said reactive functional group.